

WEST Search History

DATE: Monday, August 18, 2003

<u>Set Name</u>	<u>Query</u>	<u>Hit Count</u>	<u>Set Name</u> result set
side by side			
<i>DB=USPT; PLUR=YES; OP=AND</i>			
L1	igy.clm. and campylobact\$.clm.	0	L1
L2	ig-y.clm. and campylobact\$.clm.	0	L2
L3	campylobact\$.clm. or jejuni.clm.	191	L3
L4	L3 same (avian or turkey or fowl or poultry or chick or chicken or bird)	7	L4
L5	anticampylobacter.clm. or anti-campylobacter.clm.	0	L5
L6	anti-Campylobacter.clm.	3	L6
L7	(campylobacter or jejuni) same (igy or igg or igm or iga or antisera or antiserum or anti-sera or anti-serum or immune or immunoglobulin or monoclonal or polyclonal or poly-clonal or mono-clonal)	123	L7
L8	L7 same (avian or turkey or fowl or poultry or chick or chicken or bird)	16	L8
L9	campylobact\$.clm. or jejuni.clm.	191	L9
L10	L9 and (flge or flg-e or hook).clm.	0	L10
L11	(campylobact\$ or jejuni).ti,ab. and hook.clm.	0	L11
L12	(campylobact\$ or jejuni).ti,ab. and flgE.clm.	0	L12
L13	(campylobact\$ or jejuni) and flgE	2	L13

END OF SEARCH HISTORY

S4 1 (S1 OR S2 OR S3) AND (AVIAN (5N) (ANTIBOD? OR ANTISER? -
IMMUNE? OR SEROTYP? OR IMMUNE? OR IMMUNOGLOB? OR IGA OR IGM OR
IGG?))

S5 12196 (AVIAN OR CHICK? OR FOWL? OR TURKEY? OR BIRD? OR IGY?) (5N)
(ANTIBOD? OR ANTISER? OR IMMUNE? OR SEROTYP? OR IMMUNE? OR I-
MMUNOGLOB? OR IGA OR IGM OR IGG?)

S6 45 (S1 OR S2 OR S3) AND S5

S7 44 S6 NOT S4

S8 6 S7/2000:2003

S9 38 S7 NOT S8

?s (s1 or s2 or s3)

8257 S1

4441 S2

779 S3

S10 8313 (S1 OR S2 OR S3)

?s s10 and (western? or immunoblot? or blot?)

8313 S10

124711 WESTERN?

53434 IMMUNOBLOT?

170904 BLOT?

S11 375 S10 AND (WESTERN? OR IMMUNOBLOT? OR BLOT?)

?s s11 and (kda or kd or dalton? or kilodalton? or rmw or weight?)

375 S11

90595 KDA

44622 KD

12846 DALTON?

6642 KILODALTON?

24 RMW

550619 WEIGHT?

S12 140 S11 AND (KDA OR KD OR DALTON? OR KILODALTON? OR RMW OR
WEIGHT?)

?s s12 and (ANTIBOD? OR ANTISER? OR -

>>>Unmatched parentheses

? IMMUNE? OR SEROTYP? OR IMMUNE? OR IMMUNOGLOB? OR IGA OR IGM OR

>>>Unrecognizable Command

? IGG?))

>>>Unrecognizable Command

? s s12 and (ANTIBOD? OR ANTISER? OR igg or iga or igm or MMUNE? OR SEROTYP? OR IMMUNE? O
R IMMUNOGLOB?)

140 S12

625653 ANTIBOD?

53949 ANTISER?

76481 IGG

29714 IGA

39527 IGM

1 MMUNE?

31174 SEROTYP?

237721 IMMUNE?

205575 IMMUNOGLOB?

S13 110 S12 AND (ANTIBOD? OR ANTISER? OR IGG OR IGA OR IGM OR
MMUNE? OR SEROTYP? OR IMMUNE? OR IMMUNOGLOB?)

?s s13 and (avian? or fowl? or poultry? or turkey? or chick? or hen? or yolk? or igy or b
ird?)

110 S13

26408 AVIAN?

5994 FOWL?

21203 POULTRY?

15337 TURKEY?

127853 CHICK?

72384 HEN?

10851 YOLK?

250 IGY

34792 BIRD?

S14 9 S13 AND (AVIAN? OR FOWL? OR POULTRY? OR TURKEY? OR CHICK?
OR HEN? OR YOLK? OR IGY OR BIRD?)

?t s14/9/all

D1 MAG
8/03
108

WEST

Generate Collection

Print

Apr 12, 1994

L4: Entry 5 of 7

File: USPT

DOCUMENT-IDENTIFIER: US 5302388 A

TITLE: Control of campylobacter jejuni colonization

CLAIMS:

1. An anti-Campylobacter jejuni colonizing poultry feed which is useful for preventing the colonization of Campylobacter jejuni in a poultry animal, having dispersed therein, as an active ingredient, an effective amount of at least one cecal-colonizing strain of microorganism to provide anti-Campylobacter activity, wherein the strain is selected from the group consisting of Klebsiella pneumoniae strain 23 (ATCC No. 55234), Citrobacter diversus strain 22 (ATCC No. 55236), Escherichia coli (O13:H.sup.-) strain 25 (ATCC No. 55235), mutants thereof which retain the ability to produce anti-Campylobacter activity, and mixtures thereof.
7. The composition of claim 5, wherein the cecal-colonizing strain is an active ingredient in a poultry feed material, said strain producing anti-Campylobacter metabolites to provide, upon addition to a poultry feed, a poultry feed producing an effective amount of anti-Campylobacter metabolites to inhibit the colonization of Campylobacter jejuni in a poultry animal.
11. A process for inhibiting the colonization of Campylobacter jejuni in poultry comprising administering an effective amount of at least one cecal-colonizing strain of microorganism, said strain producing anti-Campylobacter metabolites, wherein the strain is selected from the group consisting of Klebsiella pneumoniae strain 23 (ATCC No. 55234), Citrobacter diversus strain 22 (ATCC No. 55236), Escherichia coli (O13:H.sup.-) strain 25 (ATCC No. 55235), mutants thereof which retain the ability to produce anti-Campylobacter activity, and mixtures thereof, and a carrier.
20. A process for inhibiting the colonization of Campylobacter jejuni in poultry comprising dispensing and delivering a dietary supplement comprising an effective amount of at least one cecal-colonizing strain of microorganism, said strain producing anti-Campylobacter metabolites, wherein the strain is selected from the group consisting of Klebsiella pneumoniae strain 23 (ATCC No. 55234), Citrobacter diversus strain 22 (ATCC No. 55236), Escherichia coli (O13:H.sup.-) strain 25 (ATCC No. 55235), mutants thereof which retain the ability to produce anti-Campylobacter activity, and mixtures thereof.

WEST

Generate Collection

Print

L8: Entry 10 of 16

File: USPT

Mar 30, 1999

DOCUMENT-IDENTIFIER: US 5888810 A

TITLE: Campylobacteri jejuni flagellin-escherichia coli LT-B fusion protein

Brief Summary Text (33):

Serum antibody response to invasive enteric pathogens is very important in protection against systemic infections. The initial immunologic response to enteric infection occurs at the level of the intestinal mucosa. Secretory immunoglobulin A (sIgA) response at the intestinal mucosa is a primary defense against enteric infections (Winsor et al. supra). Stern et al. (1990. Avian Dis., vol. 34, pp. 595-601) found that specific anti-C. jejuni antibodies diminish the ability of the bacterium to colonize the gut of 1-day-old chicks when incubated with the organism as compared with preincubation with phosphate buffered saline.

Brief Summary Text (34):

The flagella of C. jejuni are essential in the colonization of the intestine. Nonflagellated organisms are quickly cleared from the intestine. Chicken polyclonal antflagellin antibodies as well as monoclonal antflagellin antibodies have been found to prevent C. jejuni from colonizing the chickens or to increase the dose of bacteria required to colonize the chickens (Carr, unpublished). Flagellar antigens are therefore potential candidates for vaccines as well as suitable antigens for diagnostic purposes, since the flagellin protein is immunodominant during human infections.

Detailed Description Text (9):

The LT-B/fla fusion gene was under constitutive expression in X6097. The fusion protein was detected at several growth times. The best recovery, i.e. the greatest yield of the fusion protein relative to the total protein, was when cell density corresponding to OD_{sub}600 of about 0.8 was reached. The fusion protein was detected by Coomassie staining, and Western blot analyses using chicken anti-flagellin serum (FIG. 2), rabbit anti-C. jejuni serum, affinity purified rabbit anti-C. jejuni flagellin antibodies and rabbit anti-LT serum (FIG. 2). The fusion protein was not recognized by a monoclonal antibody directed against the 63 kd flagellin protein, presumably because the monoclonal antibody is directed against an epitope not present in our fusion protein since only 46% of the flaA gene is expressed. The fusion protein has a MW of 43 kd (16 for LT-B and 27 kd for the U band). The protein could not be detected from the pBEB transformed X6097 control. The LT-B/fla, fusin gene DNA sequence is presented in FIG. 3.

Detailed Description Text (28):

Western blot analyses were performed as described by Towbin et al (1979. PNAS, vol. 76, pp. 4350-4354). Blots were treated with a 1:200 dilution of chicken anti-C. jejuni serum or rabbit anti-LT before adding the secondary antibody (goat-anti rabbit IgG alkaline phosphatase conjugate, Bio-Rad), and developed with the substrate solution (5-bromo-4-chloro-3-indolyl phosphate/nitro blue tetrazolium) as described by Sambrook et al. (1989. In Molecular Cloning, A Laboratory Manual, Cold Spring Harbor, N.Y.: Cold Spring Harbor

Laboratory). Results are shown in FIG. 2.

S3 295 S2/2000:2003
S4 78 S2 NOT S3
?t s4/3,kwic/42

4/3,KWIC/42 (Item 17 from file: 349)

DIALOG(R)File 349:PCT FULLTEXT

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00452378 **Image available**

**A PORIN GENE FROM CAMPYLOBACTER JEJUNI, RELATED PRODUCTS AND USES THEREOF
GENE DE PORINE EXTRAIT DE CAMPYLOBACTER JEJUNI, PRODUITS APPARENTES ET
LEURS UTILISATIONS**

Patent Applicant/Assignee:

HER MAJESTY IN RIGHT OF CANADA as represented by THE MINISTER OF HEALTH
AND WELFARE CANADA,

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BACON David J,
RODGERS Frank,
BOLLA Jean-Michel,

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RODGERS Frank,
BOLLA Jean-Michel,

Patent and Priority Information (Country, Number, Date):

Patent: WO 9842842 A1 19981001

Application: WO 98CA272 19980325 (PCT/WO CA9800272)

Priority Application: US 9741200 19970325

Designated States: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES
FI GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD
MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US
UZ VN YU ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE
CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN ML
MR NE SN TD TG

Publication Language: English

Fulltext Word Count: 25116

**A PORIN GENE FROM CAMPYLOBACTER JEJUNI, RELATED PRODUCTS AND USES THEREOF
GENE DE PORINE EXTRAIT DE CAMPYLOBACTER JEJUNI, PRODUITS APPARENTES ET
LEURS UTILISATIONS**

Fulltext Availability:

Detailed Description

Claims

English Abstract

The invention relates to a porin gene from *Campylobacter jejuni* [SEQ ID NO:3]. The gene has been designated porA and is 1275 bp...

French Abstract

L'invention concerne un gene de porine extrait de *Campylobacter jejuni* (SEQ ID NO:3). Ce gene, denomme porA, a une longueur de 1275 bp...

Detailed Description

A PORIN GENE FROM *Campylobacter jejuni*,
RELATED PRODUCTS AND USES THEREOF

TECHNICAL FIELD

This invention relates to a porin gene from

5 *Campylobacter jejuni*, to related products and to the uses

BACKGROUND ART

In the following discussion, the...

...41). Active surveys

conducted in the United States have estimated the number
of cases of *campylobacteriosis* to be 2.5 million per year,
making it a multi-million dollar disease (39...

...by *C. jejuni* can range from watery to bloody
diarrhea (28, 39). In most cases *campylobacteriosis* is a
self-limiting disease but in the more severe cases,

File 155:MEDLINE(R) 1966-2003/Aug W3

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*File 155: Medline has been reloaded and accession numbers have changed. Please see HELP NEWS 155.

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      Set  Items  Description
      ---  ----  -
?e campylobacter
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Ref	Items	RT	Index-term
E1	1		CAMPYLOBACTER
E2	1		CAMPYLOBACTER
E3	8257	9	*CAMPYLOBACTER
E4	54		CAMPYLOBACTER --ANALYSIS --AN
E5	21		CAMPYLOBACTER --CHEMISTRY --CH
E6	529		CAMPYLOBACTER --CLASSIFICATION --CL
E7	15		CAMPYLOBACTER --CYTOLOGY --CY
E8	398		CAMPYLOBACTER --DRUG EFFECTS --DE
E9	125		CAMPYLOBACTER --ENZYMOLGY --EN
E10	317		CAMPYLOBACTER --GENETICS --GE
E11	270		CAMPYLOBACTER --GROWTH AND DEVELOPMENT --GD
E12	295		CAMPYLOBACTER --IMMUNOLOGY --IM

Enter P or PAGE for more

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Ref	Items	RT	Index-term
E13	1652		CAMPYLOBACTER --ISOLATION AND PURIFICATION --I
E14	130		CAMPYLOBACTER --METABOLISM --ME
E15	240		CAMPYLOBACTER --PATHOGENICITY --PY
E16	95		CAMPYLOBACTER --PHYSIOLOGY --PH
E17	6		CAMPYLOBACTER --RADIATION EFFECTS --RE
E18	79		CAMPYLOBACTER --ULTRASTRUCTURE --UL
E19	320	4	CAMPYLOBACTER COLI
E20	15		CAMPYLOBACTER COLI --CHEMISTRY --CH
E21	90		CAMPYLOBACTER COLI --CLASSIFICATION --CL
E22	57		CAMPYLOBACTER COLI --DRUG EFFECTS --DE
E23	12		CAMPYLOBACTER COLI --ENZYMOLGY --EN
E24	126		CAMPYLOBACTER COLI --GENETICS --GE

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Ref	Items	RT	Index-term
E25	32		CAMPYLOBACTER COLI --GROWTH AND DEVELOPMENT --
E26	24		CAMPYLOBACTER COLI --IMMUNOLOGY --IM
E27	148		CAMPYLOBACTER COLI --ISOLATION AND PURIFICATIO
E28	17		CAMPYLOBACTER COLI --METABOLISM --ME
E29	24		CAMPYLOBACTER COLI --PATHOGENICITY --PY
E30	16		CAMPYLOBACTER COLI --PHYSIOLOGY --PH
E31	8		CAMPYLOBACTER COLI --ULTRASTRUCTURE --UL
E32	1772	5	CAMPYLOBACTER FETUS
E33	35		CAMPYLOBACTER FETUS --ANALYSIS --AN
E34	14		CAMPYLOBACTER FETUS --CHEMISTRY --CH
E35	192		CAMPYLOBACTER FETUS --CLASSIFICATION --CL
E36	14		CAMPYLOBACTER FETUS --CYTOLOGY --CY

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Ref	Items	RT	Index-term
E37	180		CAMPYLOBACTER FETUS --DRUG EFFECTS --DE
E38	12		CAMPYLOBACTER FETUS --ENZYMOLGY --EN
E39	105		CAMPYLOBACTER FETUS --GENETICS --GE
E40	157		CAMPYLOBACTER FETUS --GROWTH AND DEVELOPMENT -
E41	241		CAMPYLOBACTER FETUS --IMMUNOLOGY --IM
E42	781		CAMPYLOBACTER FETUS --ISOLATION AND PURIFICATI
E43	80		CAMPYLOBACTER FETUS --METABOLISM --ME

E44	108	CAMPYLOBACTER FETUS --PATHOGENICITY --PY
E45	44	CAMPYLOBACTER FETUS --PHYSIOLOGY --PH
E46	2	CAMPYLOBACTER FETUS --RADIATION EFFECTS --RE
E47	43	CAMPYLOBACTER FETUS --ULTRASTRUCTURE --UL
E48	3925 2	CAMPYLOBACTER INFECTIONS

Enter P or PAGE for more

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Ref	Items	Index-term
E49	42	CAMPYLOBACTER INFECTIONS --BLOOD --BL
E50	5	CAMPYLOBACTER INFECTIONS --CEREBROSPINAL FLUID
?s e3-e50		
	8257	CAMPYLOBACTER
	54	CAMPYLOBACTER --ANALYSIS --AN
	21	CAMPYLOBACTER --CHEMISTRY --CH
	529	CAMPYLOBACTER --CLASSIFICATION --CL
	15	CAMPYLOBACTER --CYTOLOGY --CY
	398	CAMPYLOBACTER --DRUG EFFECTS --DE
	125	CAMPYLOBACTER --ENZYMولوجY --EN
	317	CAMPYLOBACTER --GENETICS --GE
	270	CAMPYLOBACTER --GROWTH AND DEVELOPMENT --GD
	295	CAMPYLOBACTER --IMMUNOLOGY --IM
	1652	CAMPYLOBACTER --ISOLATION AND PURIFICATION --I
	130	CAMPYLOBACTER --METABOLISM --ME
	240	CAMPYLOBACTER --PATHOGENICITY --PY
	95	CAMPYLOBACTER --PHYSIOLOGY --PH
	6	CAMPYLOBACTER --RADIATION EFFECTS --RE
	79	CAMPYLOBACTER --ULTRASTRUCTURE --UL
	320	CAMPYLOBACTER COLI
	15	CAMPYLOBACTER COLI --CHEMISTRY --CH
	90	CAMPYLOBACTER COLI --CLASSIFICATION --CL
	57	CAMPYLOBACTER COLI --DRUG EFFECTS --DE
	12	CAMPYLOBACTER COLI --ENZYMولوجY --EN
	126	CAMPYLOBACTER COLI --GENETICS --GE
	32	CAMPYLOBACTER COLI --GROWTH AND DEVELOPMENT --
	24	CAMPYLOBACTER COLI --IMMUNOLOGY --IM
	148	CAMPYLOBACTER COLI --ISOLATION AND PURIFICATIO
	17	CAMPYLOBACTER COLI --METABOLISM --ME
	24	CAMPYLOBACTER COLI --PATHOGENICITY --PY
	16	CAMPYLOBACTER COLI --PHYSIOLOGY --PH
	8	CAMPYLOBACTER COLI --ULTRASTRUCTURE --UL
	1772	CAMPYLOBACTER FETUS
	35	CAMPYLOBACTER FETUS --ANALYSIS --AN
	14	CAMPYLOBACTER FETUS --CHEMISTRY --CH
	192	CAMPYLOBACTER FETUS --CLASSIFICATION --CL
	14	CAMPYLOBACTER FETUS --CYTOLOGY --CY
	180	CAMPYLOBACTER FETUS --DRUG EFFECTS --DE
	12	CAMPYLOBACTER FETUS --ENZYMولوجY --EN
	105	CAMPYLOBACTER FETUS --GENETICS --GE
	157	CAMPYLOBACTER FETUS --GROWTH AND DEVELOPMENT -
	241	CAMPYLOBACTER FETUS --IMMUNOLOGY --IM
	781	CAMPYLOBACTER FETUS --ISOLATION AND PURIFICATI
	80	CAMPYLOBACTER FETUS --METABOLISM --ME
	108	CAMPYLOBACTER FETUS --PATHOGENICITY --PY
	44	CAMPYLOBACTER FETUS --PHYSIOLOGY --PH
	2	CAMPYLOBACTER FETUS --RADIATION EFFECTS --RE
	43	CAMPYLOBACTER FETUS --ULTRASTRUCTURE --UL
	3925	CAMPYLOBACTER INFECTIONS
	42	CAMPYLOBACTER INFECTIONS --BLOOD --BL
	5	CAMPYLOBACTER INFECTIONS --CEREBROSPINAL FLUID
S1	8257	E3-E50

?p

Ref	Items	Index-term
E1	5	CAMPYLOBACTER INFECTIONS --CEREBROSPINAL FLUID
E2	4	CAMPYLOBACTER INFECTIONS --CHEMICALLY INDUCED
E3	7	CAMPYLOBACTER INFECTIONS --CLASSIFICATION --CL
E4	713	CAMPYLOBACTER INFECTIONS --COMPLICATIONS --CO

E5	3	CAMPYLOBACTER INFECTIONS --CONGENITAL --CN
E6	614	CAMPYLOBACTER INFECTIONS --DIAGNOSIS --DI
E7	2	CAMPYLOBACTER INFECTIONS --DIET THERAPY --DH
E8	477	CAMPYLOBACTER INFECTIONS --DRUG THERAPY --DT
E9	11	CAMPYLOBACTER INFECTIONS --ECONOMICS --EC
E10	3	CAMPYLOBACTER INFECTIONS --ENZYMولوجY --EN
E11	906	CAMPYLOBACTER INFECTIONS --EPIDEMIOLOGY --EP
E12	3	CAMPYLOBACTER INFECTIONS --ETHNOLOGY --EH

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Ref	Items	Index-term
E13	268	CAMPYLOBACTER INFECTIONS --ETIOLOGY --ET
E14	23	CAMPYLOBACTER INFECTIONS --GENETICS --GE
E15	6	CAMPYLOBACTER INFECTIONS --HISTORY --HI
E16	335	CAMPYLOBACTER INFECTIONS --IMMUNOLOGY --IM
E17	27	CAMPYLOBACTER INFECTIONS --METABOLISM --ME
E18	1281	CAMPYLOBACTER INFECTIONS --MICROBIOLOGY --MI
E19	13	CAMPYLOBACTER INFECTIONS --MORTALITY --MO
E20	2	CAMPYLOBACTER INFECTIONS --NURSING --NU
E21	249	CAMPYLOBACTER INFECTIONS --PATHOLOGY --PA
E22	78	CAMPYLOBACTER INFECTIONS --PHYSIOPATHOLOGY --P
E23	140	CAMPYLOBACTER INFECTIONS --PREVENTION AND CONT
E24	4	CAMPYLOBACTER INFECTIONS --PSYCHOLOGY --PX

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Ref	Items	RT	Index-term
E25	10		CAMPYLOBACTER INFECTIONS --RADIOGRAPHY --RA
E26	11		CAMPYLOBACTER INFECTIONS --SURGERY --SU
E27	61		CAMPYLOBACTER INFECTIONS --THERAPY --TH
E28	299		CAMPYLOBACTER INFECTIONS --TRANSMISSION --TM
E29	3		CAMPYLOBACTER INFECTIONS --ULTRASONOGRAPHY --U
E30	1		CAMPYLOBACTER INFECTIONS --URINE --UR
E31	578		CAMPYLOBACTER INFECTIONS --VETERINARY --VE
E32	1577	4	CAMPYLOBACTER JEJUNI
E33	2		CAMPYLOBACTER JEJUNI --ANALYSIS --AN
E34	67		CAMPYLOBACTER JEJUNI --CHEMISTRY --CH
E35	259		CAMPYLOBACTER JEJUNI --CLASSIFICATION --CL
E36	12		CAMPYLOBACTER JEJUNI --CYTOLOGY --CY

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Ref	Items	RT	Index-term
E37	190		CAMPYLOBACTER JEJUNI --DRUG EFFECTS --DE
E38	39		CAMPYLOBACTER JEJUNI --ENZYMولوجY --EN
E39	425		CAMPYLOBACTER JEJUNI --GENETICS --GE
E40	155		CAMPYLOBACTER JEJUNI --GROWTH AND DEVELOPMENT
E41	243		CAMPYLOBACTER JEJUNI --IMMUNOLOGY --IM
E42	553		CAMPYLOBACTER JEJUNI --ISOLATION AND PURIFICAT
E43	81		CAMPYLOBACTER JEJUNI --METABOLISM --ME
E44	179		CAMPYLOBACTER JEJUNI --PATHOGENICITY --PY
E45	80		CAMPYLOBACTER JEJUNI --PHYSIOLOGY --PH
E46	1		CAMPYLOBACTER JEJUNI --RADIATION EFFECTS --RE
E47	36		CAMPYLOBACTER JEJUNI --ULTRASTRUCTURE --UL
E48	0	1	CAMPYLOBACTER PYLORI

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?s e1-e47

5	CAMPYLOBACTER INFECTIONS --CEREBROSPINAL FLUID
4	CAMPYLOBACTER INFECTIONS --CHEMICALLY INDUCED
7	CAMPYLOBACTER INFECTIONS --CLASSIFICATION --CL
713	CAMPYLOBACTER INFECTIONS --COMPLICATIONS --CO
3	CAMPYLOBACTER INFECTIONS --CONGENITAL --CN
614	CAMPYLOBACTER INFECTIONS --DIAGNOSIS --DI
2	CAMPYLOBACTER INFECTIONS --DIET THERAPY --DH

477	CAMPYLOBACTER INFECTIONS	--DRUG THERAPY --DT
11	CAMPYLOBACTER INFECTIONS	--ECONOMICS --EC
3	CAMPYLOBACTER INFECTIONS	--ENZYMولوجY --EN
906	CAMPYLOBACTER INFECTIONS	--EPIDEMIOLOGY --EP
3	CAMPYLOBACTER INFECTIONS	--ETHNOLOGY --EH
268	CAMPYLOBACTER INFECTIONS	--ETIOLOGY --ET
23	CAMPYLOBACTER INFECTIONS	--GENETICS --GE
6	CAMPYLOBACTER INFECTIONS	--HISTORY --HI
335	CAMPYLOBACTER INFECTIONS	--IMMUNOLOGY --IM
27	CAMPYLOBACTER INFECTIONS	--METABOLISM --ME
1281	CAMPYLOBACTER INFECTIONS	--MICROBIOLOGY --MI
13	CAMPYLOBACTER INFECTIONS	--MORTALITY --MO
2	CAMPYLOBACTER INFECTIONS	--NURSING --NU
249	CAMPYLOBACTER INFECTIONS	--PATHOLOGY --PA
78	CAMPYLOBACTER INFECTIONS	--PHYSIOPATHOLOGY --P
140	CAMPYLOBACTER INFECTIONS	--PREVENTION AND CONT
4	CAMPYLOBACTER INFECTIONS	--PSYCHOLOGY --PX
10	CAMPYLOBACTER INFECTIONS	--RADIOGRAPHY --RA
11	CAMPYLOBACTER INFECTIONS	--SURGERY --SU
61	CAMPYLOBACTER INFECTIONS	--THERAPY --TH
299	CAMPYLOBACTER INFECTIONS	--TRANSMISSION --TM
3	CAMPYLOBACTER INFECTIONS	--ULTRASONOGRAPHY --U
1	CAMPYLOBACTER INFECTIONS	--URINE --UR
578	CAMPYLOBACTER INFECTIONS	--VETERINARY --VE
1577	CAMPYLOBACTER JEJUNI	
2	CAMPYLOBACTER JEJUNI	--ANALYSIS --AN
67	CAMPYLOBACTER JEJUNI	--CHEMISTRY --CH
259	CAMPYLOBACTER JEJUNI	--CLASSIFICATION --CL
12	CAMPYLOBACTER JEJUNI	--CYTOLOGY --CY
190	CAMPYLOBACTER JEJUNI	--DRUG EFFECTS --DE
39	CAMPYLOBACTER JEJUNI	--ENZYMولوجY --EN
425	CAMPYLOBACTER JEJUNI	--GENETICS --GE
155	CAMPYLOBACTER JEJUNI	--GROWTH AND DEVELOPMENT
243	CAMPYLOBACTER JEJUNI	--IMMUNOLOGY --IM
553	CAMPYLOBACTER JEJUNI	--ISOLATION AND PURIFICAT
81	CAMPYLOBACTER JEJUNI	--METABOLISM --ME
179	CAMPYLOBACTER JEJUNI	--PATHOGENICITY --PY
80	CAMPYLOBACTER JEJUNI	--PHYSIOLOGY --PH
1	CAMPYLOBACTER JEJUNI	--RADIATION EFFECTS --RE
36	CAMPYLOBACTER JEJUNI	--ULTRASTRUCTURE --UL
S2	4441	E1-E47

?p

Ref	Items	Index-term
E49	3	CAMPYLOBACTERACEAE
E50	1	CAMPYLOBACTERAHNNLICHE

?p

Ref	Items	Index-term
E1	1	CAMPYLOBACTERAHNNLICHE
E2	1	CAMPYLOBACTERARTEN
E3	1	CAMPYLOBACTERCINAEDI
E4	1	CAMPYLOBACTERENTERIT
E5	1	CAMPYLOBACTEREPIDEMIOLOGI
E6	37	CAMPYLOBACTERIA
E7	1	CAMPYLOBACTERIACEAE
E8	1	CAMPYLOBACTERIAL
E9	1	CAMPYLOBACTERIASIS
E10	1	CAMPYLOBACTERIES
E11	2	CAMPYLOBACTERINFEKSJON
E12	1	CAMPYLOBACTERINFEKTION

Enter P or PAGE for more

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Ref	Items	Index-term
E13	3	CAMPYLOBACTERINFEKTIONEN
E14	8	CAMPYLOBACTERIOSE
E15	1	CAMPYLOBACTERIOSES

E16	266	CAMPYLOBACTERIOSIS
E17	2	CAMPYLOBACTERIUM
E18	9	CAMPYLOBACTERJEJUNI
E19	2	CAMPYLOBACTERLIKE
E20	1	CAMPYLOBACTERN
E21	6	CAMPYLOBACTEROSIS
E22	462	CAMPYLOBACTERS
E23	1	CAMPYLOBACTERSPECIES
E24	2	CAMPYLOBACTERSPP

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Ref	Items	Index-term
E25	1	CAMPYLOBACTERSTAMME
E26	1	CAMPYLOBACTERY
E27	2	CAMPYLOBACTOR
E28	1	CAMPYLOBACTWER
E29	1	CAMPYLOBACYTER
E30	1	CAMPYLOBAKTER
E31	1	CAMPYLOBAKTERIOSE
E32	2	CAMPYLOBAKTEROVE
E33	1	CAMPYLOBASTER
E34	3	CAMPYLOBATER
E35	1	CAMPYLOBATERIOSES
E36	1	CAMPYLOBATERS

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Ref	Items	Index-term
E37	1	CAMPYLOBYCTER
E38	1	CAMPYLODACTYLI
E39	1	CAMPYLODACTYLIA
E40	1	CAMPYLODACTYLIE
E41	1	CAMPYLODACTYLY
E42	1	CAMPYLOGNATHIA
E43	1	CAMPYLOMMA
E44	2	CAMPYLOMORMYRUS
E45	1	CAMPYLOPODA
E46	1	CAMPYLOPODIUM
E47	1	CAMPYLOPTERUS
E48	1	CAMPYLOPUS

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?s e1-ee37

>>>Invalid term: EE37

?ds

Set	Items	Description
S1	8257	E3-E50
S2	4441	E1-E47

?s e1-e37

1	CAMPYLOBACTERAHNLCHE
1	CAMPYLOBACTERARTEN
1	CAMPYLOBACTERCINAEDI
1	CAMPYLOBACTERENTERIT
1	CAMPYLOBACTEREPIDEMIOLOGI
37	CAMPYLOBACTERIA
1	CAMPYLOBACTERIACEAE
1	CAMPYLOBACTERIAL
1	CAMPYLOBACTERIASIS
1	CAMPYLOBACTERIES
2	CAMPYLOBACTERINFEKSJON
1	CAMPYLOBACTERINFEKTION
3	CAMPYLOBACTERINFEKTIONEN
8	CAMPYLOBACTERIOSE
1	CAMPYLOBACTERIOSES
266	CAMPYLOBACTERIOSIS

2 CAMPYLOBACTERIUM
 9 CAMPYLOBACTERJEJUNI
 2 CAMPYLOBACTERLIKE
 1 CAMPYLOBACTERN
 6 CAMPYLOBACTEROSIS
 462 CAMPYLOBACTERS
 1 CAMPYLOBACTERSPECIES
 2 CAMPYLOBACTERSPP
 1 CAMPYLOBACTERSTAMME
 1 CAMPYLOBACTERY
 2 CAMPYLOBACTOR
 1 CAMPYLOBACTWER
 1 CAMPYLOBACYTER
 1 CAMPYLOBAKTER
 1 CAMPYLOBAKTERIOSE
 2 CAMPYLOBAKTEROVE
 1 CAMPYLOBASTER
 3 CAMPYLOBATER
 1 CAMPYLOBATERIOSES
 1 CAMPYLOBATERS
 1 CAMPYLOBYCTER

S3 779 E1-E37

?ds

Set	Items	Description
S1	8257	E3-E50
S2	4441	E1-E47
S3	779	E1-E37

?s (s1 or s2 or s3) and (avian (5n) (antibod? or antiser? or immune? or serotyp? or immun
e? or immunoglob? or iga or igm or igg?))

8257 S1
 4441 S2
 779 S3
 26246 AVIAN
 625653 ANTIBOD?
 53949 ANTISER?
 237721 IMMUNE?
 31174 SEROTYP?
 237721 IMMUNE?
 205575 IMMUNOGLOB?
 29714 IGA
 39527 IGM
 85819 IGG?
 1171 AVIAN(5N)((((((((ANTIBOD? OR ANTISER?) OR IMMUNE?) OR
 SEROTYP?) OR IMMUNE?) OR IMMUNOGLOB?) OR IGA) OR IGM) OR
 IGG?)

S4 1 (S1 OR S2 OR S3) AND (AVIAN (5N) (ANTIBOD? OR ANTISER? OR
IMMUNE? OR SEROTYP? OR IMMUNE? OR IMMUNOGLOB? OR IGA OR
IGM OR IGG?))

?t s4/9/all

4/9/1

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05507901 87186825 PMID: 3567742

A study of factors affecting the sensitivity of the passive haemagglutination method for serotyping Campylobacter jejuni and Campylobacter coli and recommendations for a more rapid procedure.

Fricker C R; Alemohammad M M; Park R W

Canadian journal of microbiology (CANADA) Jan 1987, 33 (1) p33-9,
ISSN 0008-4166 Journal Code: 0372707

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Factors affecting the sensitivity of the passive haemagglutination method for serotyping campylobacters have been studied. The concentration of red

blood cells during the haemagglutination stage of the procedure markedly affected the titer obtained. An increase in concentration of red blood cells resulted in a lower titer, with titers being inversely proportional to red blood cell concentration. No differences in titer were observed when erythrocytes were sensitized at a range of pH values between pH 5.0 and pH 8.0. The time required for antigen extraction and for red blood cell sensitization was shown to be 15 min each, thus resulting in a reduction in the time required for **serotyping**. Furthermore, use of **avian** erythrocytes enabled the haemagglutination reactions to be read after incubation for only 1 h. Combining these procedures with a rapid slide haemagglutination test enables a single worker to serotype over 100 C. jejuni and C. coli isolates within 1 working day.

Tags: Animal; Comparative Study; Human; Support, Non-U.S. Gov't

Descriptors: **Campylobacter** --classification--CL; * **Campylobacter** fetus --classification--CL; *Hemagglutination Tests--methods--MT; Antigens, Bacterial; Chickens--blood--BL; Hydrogen-Ion Concentration; Serotyping --methods--MT; Sheep--blood--BL; Time Factors; Turkeys--blood--BL

CAS Registry No.: 0 (Antigens, Bacterial)

Record Date Created: 19870526

Record Date Completed: 19870526

?s(avian or chick? or fowl? or turkey? or bird? or igy?) (5n) (antibod? or antiser? or immune? or serotyp? or immune? or immunoglob? or iga or igm or igg?)

26246 AVIAN
127853 CHICK?
5994 FOWL?
15337 TURKEY?
34792 BIRD?
250 IGY?
625653 ANTIBOD?
53949 ANTISER?
237721 IMMUNE?
31174 SEROTYP?
237721 IMMUNE?
205575 IMMUNOGLOB?
29714 IGA
39527 IGM
85819 IGG?
S5 12196 (AVIAN OR CHICK? OR FOWL? OR TURKEY? OR BIRD? OR IGY?)
(5N) (ANTIBOD? OR ANTISER? OR IMMUNE? OR SEROTYP? OR
IMMUNE? OR IMMUNOGLOB? OR IGA OR IGM OR IGG?)

?ds

Set	Items	Description
S1	8257	E3-E50
S2	4441	E1-E47
S3	779	E1-E37
S4	1	(S1 OR S2 OR S3) AND (AVIAN (5N) (ANTIBOD? OR ANTISER? OR - IMMUNE? OR SEROTYP? OR IMMUNE? OR IMMUNOGLOB? OR IGA OR IGM OR IGG?))
S5	12196	(AVIAN OR CHICK? OR FOWL? OR TURKEY? OR BIRD? OR IGY?) (5N) (ANTIBOD? OR ANTISER? OR IMMUNE? OR SEROTYP? OR IMMUNE? OR I- MMUNOGLOB? OR IGA OR IGM OR IGG?)

?s (s1 or s2 or s3) and s5

8257 S1
4441 S2
779 S3
12196 S5

S6 45 (S1 OR S2 OR S3) AND S5

?s s6 not s4

45 S6
1 S4

S7 44 S6 NOT S4

?s s7/2000:2003

44 S7
1844888 PY=2000 : PY=2003

S8 6 S7/2000:2003

?s s7 not s8

44 S7
6 S8

S9 38 S7 NOT S8
?t s9/9/all

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Set Items Description

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Set Items Description

S1 7276 (AVIAN? OR CHICKEN? OR TURKEY? OR FOWL? OR BIRD? OR IGY?) -
(100N) (ANTIBOD? OR MONOCLON? OR POLYCLONAL? OR ANTISER? OR I-
MMUNE? OR IMMUNOGLOB?) (100N) (FLAGELLALESS? OR NONFLAGEL? OR
AFLAGEL? OR KDA OR K)
S2 373 S1 AND CAMPYLOBAC?

antibiotic intervention with macrolids...

...34). Johnson

5 and Lior (19) originally reported that 410 of 718 isolates of *Campylobacter* sp. screened for the production of CLDT were positive; however, isolates screened for the cdt13...said target.

The invention also relates to a method of detecting the presence of *Campylobacter* jejuni infection, characterized by the steps of: a) contacting a sample obtained from a patient...

...a time sufficient to allow formation of a complex between said protein and any anti- *Campylobacter* jejuni antibodies present in said sample; and b) detecting the presence of, and optionally the...

...comprises an isolated expression vector, characterized by a region encoding a porA protein of *Campylobacter* jejuni, or an antigenic fragment thereof.

Included within the invention is a method of inducing...

...aspect of the invention is a method of producing antibodies for testing for infection by *Campylobacter* jejuni, characterized in that a protein having an amino acid sequence of SEQ ID NO...h; Figure 8 shows western blot analysis of the isolated cytotoxic porin-LPS complex from *Campylobacter* sp using 40 Ag of crude, concentrated filtrate and homologous rabbit antiserum. Lanes 1 and...present invention is based on the identification of a porin-lipopolysaccharide (LPS) complex from *Campylobacter* jejuni that is an endotoxin and that is fairly well conserved amongst strains of the organism, but not widely found in other *Campylobacter* species. The complex has been isolated and a corresponding porin gene, designated 11porA, 11...subsequently stored at -700C in tryptic soy broth containing 5% sheep blood. Strains of *Campylobacter* sp.

and related organisms were maintained at -800C in glycerol-peptone water as part of...ompC of *S. typhi* (29), a 45i similarity and 200i identity was found. Screening *Campylobacter* sp. for porA and cytotoxin production.

Results of screening *C. jejuni* for phenotypic ...gene are summarized in Table 2. It was found that all 32 strains of *Campylobacter* sp. and related organisms produced a cytotoxic component when the filtrate from the biphasic -44...

...of *C. jejuni*, especially Lior serotype 82, but was not conserved between related species of *Campylobacter*.

DISCUSSION

The 1275 bp ORF had a '%guanosine+cytosine content of 36.8 mol % (Fig...contained at least

part, if not all of the intact gene while the other *Campylobacter* sp. and related organism were PCR negative.

Previous reports indicated that only 60% of...are valuable, and provide a new and efficient method to identify *C. jejuni* from other

Campylobacter sp. The potential for the development of a recombinant vaccine using the porin protein is...on

immuno-blot analysis. Western blots of crude concentrated filtrates from various cytotoxic strains of

Campylobacter species showed the presence of a protein with a molecular mass similar to that of...comparable molecular weight was also present in crude concentrated filtrates from other cytotoxic strains of **Campylobacter** sp., indicating that the release of the porin-LPS complex 25 was not unique to...been cloned and sequenced, a fuller understanding of the role of the porin in clinical **campylobacteriosis** will be forthcoming. Such evaluations may suggest potential roles for the porin-LPS complex as...COUNTRY: France

(F) POSTAL CODE (ZIP): 13009

(ii) TITLE OF INVENTION: A PORIN GENE FROM **CAMPYLOBACTER** JEJUNI, RELATED PRODUCTS AND USES THEREOF

(iii) NUMBER OF SEQUENCES: 31

-70 (iv) COMPUTER READABLE...unknown

(ii) MOLECULE TYPE: protein

(v) FRAGMENT TYPE: N-terminal

(vi) ORIGINAL SOURCE.

(A) ORGANISM: **Campylobacter jejuni**

(B) STRAIN: 2483

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2.

Met Lys Leu Val...STRANDEDNESS: double

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(vi) ORIGINAL SOURCE.

(A) ORGANISM: **Campylobacter jejuni**

(B) STRAIN: 2483

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3.

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Claim

... purified nucleic acid, characterized in that said nucleic acid encodes a porA protein of **Campylobacter jejuni**, or an antigenic fragment thereof.

2 A nucleic acid according to claim 1, characterized...

...acid according to claim 1, characterized in that it is derived from strain 2483 of **Campylobacter** 10 jejuni (ATCC Accession No.

4 A nucleic acid according to claim 1, characterized in...

...probe to bind specifically to said target.

59 A method of detecting the presence of **Campylobacter jejuni** infection, characterized by the steps of:
a) contacting a sample obtained from a patient...

...time sufficient to
10 allow formation of a complex between said protein and any anti- **Campylobacter jejuni** antibodies present in said sample; and
b) detecting the presence of, and optionally the...

...said complex formed during step (a).
15 10. A method of detecting the presence of **Campylobacter jejuni** in a patient, characterized by obtaining from said patient a sample suspected of containing **Campylobacter jejuni**, and detecting whether the characteristic nucleic acid of claim 1, claim 2, claim 3...

...103

14 An isolated expression vector, characterized by a region encoding a porA protein of **Campylobacter jejuni**, or an antigenic ...said complex.

19 A vaccine comprising an immunogenically effective amount of the porA antigen of **Campylobacter** ' ' . or 3e3un7

antigenic fragment thereof and a pharmaceutically acceptable carrier.

20 20. A vaccine, characterized...

...human or animal body.

22 A method of producing antibodies for testing for infection by **Campylobacter jejuni**, characterized in that a 5 protein having an amino acid sequence of SEQ ID...

?t s4/9/62 64 68

4/9/62 (Item 3 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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10109393 BIOSIS NO.: 199698564311

Identification and characterization of an immunogenic outer membrane protein of *Campylobacter jejuni*.

AUTHOR: Burnens Andre; Stucki Urs; Nicolet Jacques; Frey Joachim(a)

AUTHOR ADDRESS: (a)Inst. Veterinary Bacteriol., Univ. Berne,
Langgassstrasse 122, CH-3012 Berne**Switzerland

JOURNAL: Journal of Clinical Microbiology 33 (11):p2826-2832 1995

ISSN: 0095-1137

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: We cloned and expressed in *Escherichia coli* a gene encoding an 18- kDa outer membrane protein (Omp18) from *Campylobacter jejuni* ATCC 29428. The nucleotide sequence of the gene encoding Omp18 was determined, and an open reading frame of 165 amino acids was revealed. The amino acid sequence had the typical features of a leader sequence and a signal peptidase II cleavage site at the N-terminal part of Omp18. Moreover, the sequence had a high degree of similarity to the peptidoglycan-associated outer membrane lipoprotein P6 of *Haemophilus influenzae* and the peptidoglycan-associated lipoprotein PAL of *E. coli*. Southern blot analysis in which the cloned gene was used as a probe revealed genes similar to that encoding Omp18 in all species of the thermophilic group of *campylobacters* as well as *Campylobacter sputorum*. All *campylobacters* tested expressed a protein with a molecular mass identical to that of Omp18. The protein reacted immunologically with polyclonal antibodies directed against Omp18 from *C. jejuni*. PCR amplification of the gene encoding Omp18 with specific primers and subsequent restriction enzyme analysis of the amplified DNA fragments showed that the gene for Omp18 is highly conserved in *C. jejuni* strains isolated from humans, dogs, cats, calves, and chickens but is different in other *Campylobacter* species. In order to obtain pure recombinant Omp18 protein for serological assays, the cloned gene for Omp18 was genetically modified by replacing the signal sequence with a DNA segment encoding six adjacent histidine residues. Expression of this construct in *E. coli* allowed showed no reaction with this antigen. Omp18, which is an outer membrane protein belonging to the family of PALs is well conserved in *C. jejuni* and is highly immunogenic. It is therefore a good candidate as an antigen for the serological diagnosis of past *C. jejuni* infections.

REGISTRY NUMBERS: 170613-04-4: GENBANK-X83374

DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Blood and Lymphatics (Transport and Circulation); Immune System (Chemical Coordination and Homeostasis); Infection; Membranes (Cell Biology); Pathology; Veterinary Medicine (Medical Sciences)

BIOSYSTEMATIC NAMES: Aerobic Helical or Vibrioid Gram-Negatives--Eubacteria, Bacteria; Bovidae--Artiodactyla, Mammalia, Vertebrata, Chordata, Animalia; Canidae--Carnivora, Mammalia, Vertebrata, Chordata, Animalia; Felidae--Carnivora, Mammalia, Vertebrata, Chordata, Animalia; Galliformes--Aves, Vertebrata, Chordata, Animalia; Hominidae--Primates, Mammalia, Vertebrata, Chordata, Animalia; Pasteurellaceae--Eubacteria, Bacteria; Trogoniformes--Aves, Vertebrata, Chordata, Animalia

ORGANISMS: aerobic helical or vibrioid gram-negative bacteria (Aerobic Helical or Vibrioid Gram-Negatives); calf (Bovidae); cat (Felidae); chicken (Galliformes); dog (Canidae); human (Hominidae); *Campylobacter jejuni* (Aerobic Helical or Vibrioid Gram-Negatives); *Campylobacter sputorum* (Aerobic Helical or Vibrioid Gram-Negatives); *Haemophilus influenzae* (Pasteurellaceae); Trogoniformes (Trogoniformes)

BIOSYSTEMATIC CLASSIFICATION (SUPER TAXA): animals; artiodactyls; bacteria; birds; carnivores; chordates; eubacteria; humans; mammals; microorganisms; nonhuman mammals; nonhuman vertebrates; primates; vertebrates

CHEMICALS & BIOCHEMICALS: GENBANK-X83374

MOLECULAR SEQUENCE DATABANK NUMBER: molecular sequence data; nucleotide sequence; EMBL-X83374; GENBANK-X83374

MISCELLANEOUS TERMS: INFECTION; OPEN READING FRAME; OUTER MEMBRANE PROTEIN 18 GENE; PEPTIDOGLYCAN-ASSOCIATED OUTER MEMBRANE LIPOPROTEIN P6 SIMILARITY; SEROLOGICAL DIAGNOSIS

CONCEPT CODES:

\$2.54 Estimated cost File349
 \$0.30 0.054 DialUnits File5
 \$3.50 2 Type(s) in Format 9
 \$3.50 2 Types
 \$3.80 Estimated cost File5
 \$0.04 0.009 DialUnits File35
 \$0.04 Estimated cost File35
 \$0.05 0.018 DialUnits File10
 \$1.35 1 Type(s) in Format 9
 \$1.35 1 Types
 \$1.40 Estimated cost File10
 \$0.03 0.009 DialUnits File144
 \$0.03 Estimated cost File144
 \$0.10 0.009 DialUnits File347
 \$0.10 Estimated cost File347
 \$0.05 0.009 DialUnits File160
 \$0.05 Estimated cost File160
 \$0.03 0.009 DialUnits File65
 \$0.03 Estimated cost File65
 \$0.02 0.009 DialUnits File143
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 \$0.14 0.009 DialUnits File340
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 \$0.02 0.009 DialUnits File203
 \$0.02 Estimated cost File203
 \$0.06 0.009 DialUnits File185
 \$0.06 Estimated cost File185
 \$0.03 0.009 DialUnits File94
 \$0.03 Estimated cost File94
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 \$0.05 Estimated cost File636
 \$0.04 0.009 DialUnits File444
 \$0.04 Estimated cost File444
 \$0.05 0.009 DialUnits File16
 \$0.05 Estimated cost File16
 OneSearch, 21 files, 0.430 DialUnits FileOS
 \$0.22 TELNET
 \$8.85 Estimated cost this search
 \$8.85 Estimated total session cost 0.430 DialUnits

Status: Signed Off. (1 minutes)

10973131 97325879 PMID: 9182891

Oral administration of antibodies as prophylaxis and therapy in
Campylobacter jejuni-infected chickens.

Tsubokura K; Berndtson E; Bogstedt A; Kaijser B; Kim M; Ozeki M;
Hammarstrom L

Department of Clinical Immunology, Huddinge Hospital, Sweden.

Clinical and experimental immunology (ENGLAND) Jun 1997, 108 (3)
p451-5, ISSN 0009-9104 Journal Code: 0057202

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

Passive immunity against gastrointestinal infections has recently been successfully applied as prophylaxis and therapy in patients in a variety of virally and bacterially induced infections. *Campylobacter jejuni* is frequently associated with acute diarrhoea in humans, and several species of animals have been shown to transmit the disease, although birds have been implicated as the main source of infection. We used bovine and chicken immunoglobulin preparations from the milk and eggs, respectively, of immunized animals for prophylactic and therapeutic treatment of chickens infected with *C. jejuni*. A marked prophylactic effect (a >99% decrease in the number of bacteria) was noted using either antibody preparation, whereas the therapeutic efficacy, i.e. when antibodies were given after the infection was established, was distinctly lower (80-95%) as judged by faecal bacterial counts. These observations may serve as a starting point for experiments aimed at elimination of the infection in an industrial or farm setting. It may also encourage future attempts to treat, prophylactically or therapeutically, patients with *Campylobacter*-induced diarrhoea.

Tags: Animal; Female; Support, Non-U.S. Gov't

Descriptors: Antibodies, Bacterial--therapeutic use--TU; * *Campylobacter*
Infections--prevention and control--PC; * *Campylobacter jejuni*
--immunology--IM; Administration, Oral; *Campylobacter* Infections--therapy
--TH; Cattle; Chickens; Immunization, Passive

CAS Registry No.: 0 (Antibodies, Bacterial)

Record Date Created: 19970626

Record Date Completed: 19970626

11654827 99089498 PMID: 9874102

The specificity of antibody in chickens immunised to reduce intestinal colonisation with Campylobacter jejuni.

Widders P R; Thomas L M; Long K A; Tokhi M A; Panaccio M; Apos E
Australian Quarantine and Inspection Service, Mascot, NSW.
phillip.widders@dpi.gov.au

Veterinary microbiology (NETHERLANDS) Nov 1998, 64 (1) p39-50,
ISSN 0378-1135 Journal Code: 7705469

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

Poultry consumption has been identified as a major risk factor for human infection with **Campylobacter jejuni** in developed countries. **C. jejuni** is present in the gastrointestinal tract of broiler **chickens** at the time of slaughter, and faecal contamination of carcasses during processing results in significant **campylobacter** loads on carcasses. One approach to reducing the level of carcass contamination with **C. jejuni** is to control **campylobacter** infection in broiler **chickens**. To this end, the study described here investigated the specificity of **antibody** in serum and intestinal secretions of **chickens** that had been immunised with **campylobacter** antigens and then challenged with viable bacteria. The immunodominant antigens in the serum of **birds** that showed a 2-log reduction in caecal colonisation with **C. jejuni** included flagellin protein (61-63 Kd) and three additional antigens of 67, 73.5 and 77.5 Kd. Only flagellin and the 67 Kd antigen were recognised by **IgG antibody** in gastrointestinal secretions of the same **birds**. **Antibody** from **chickens** immunised with purified native flagellin protein recognised flagellin protein and the 67 Kd antigen in **Western blots** probed with serum, but only the flagellin proteins (61-63 Kd) in **Westerns** probed with gastrointestinal secretions. Analysis of the specificity of the response to flagellin protein using recombinant clones that expressed regions of the flagellin gene suggests that epitopes in each region of the flagellin protein were immunogenic. Of the immunodominant antigens, only flagellin appeared to be surface-exposed on viable **C. jejuni**, although conformational epitopes of flagellin appeared to be sensitive to the method of antigen purification. The results of this study suggest that flagellin and possibly the 67 Kd antigen may be valuable for immunological control of intestinal infection with **C. jejuni** in **chickens**, but that further work is required to purify these as vaccine candidates by using methods that preserve conformational epitopes.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: **Antibodies**, Bacterial--immunology--IM; * **Antibody**
Specificity; * **Campylobacter** Infections--veterinary--VE; * **Campylobacter**

11829439 99269532 PMID: 10337238

Biotin-streptavidin enzyme-linked immunosorbent assay for the detection of antibodies to Campylobacter jejuni and C. coli in chickens.

Haas B; Hinz K H; Glunder G

Clinic for Poultry, Hanover School of Veterinary Medicine, Germany.

Zentralblatt fur Veterinarmedizin. Reihe B. Journal of veterinary medicine. Series B (GERMANY) Apr 1999, 46 (3) p163-71, ISSN 0514-7166
Journal Code: 0331325

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

An enzyme-linked immunosorbent assay (ELISA) was developed in a homologous system with bacterial ultrasonic-treated proteins as the antigen and antisera from chickens infected orally and subcutaneously with the strain **Campylobacter jejuni** serovar 6 (CJ 6). The cut-off level was determined using antisera from non-infected specific-pathogen-free chickens up to the age of 10 weeks. The suitability of the ELISA system was verified using antisera taken from chickens orally infected at the age of 4 weeks with CJ 1, 6, 28 or 36 or with **Campylobacter coli** serovar 28 (CC 28). The development of antibodies was monitored up to 6 weeks post-infection (p.i.). Sera from chickens infected with CJ 1, 6, 36 or CC 28 contained specific antibodies to **Campylobacter**, whereas in those infected with CJ 28 no specific antibodies were found. Distinct cross-reactions were observed between CJ 6, 28 and CC 28 antigens and their antisera 6 weeks p.i., while poor cross-reactions were found with antisera to CJ 1 and 28. Antibodies to strains of all heterologous serovars were successfully detected with an antigen pool comprised of CJ 1, 6 and 36 antigens. In 11 out of the 12 field sera obtained from 5- and 9-week-old broiler chickens suffering from **campylobacteriosis**, high specific antibody titres to **Campylobacter jejuni** were found.

Tags: Animal

10700234 '97049491 PMID: 8894221

Immunisation of chickens to reduce intestinal colonisation with Campylobacter jejuni.

Widders P R; Perry R; Muir W I; Husband A J; Long K A

Department of Agriculture, Energy and Minerals, Victorian Institute of Animal Science, Attwood, Australia.

British poultry science (ENGLAND) Sep 1996, 37 (4) p765-78, ISSN 0007-1668 Journal Code: 15740290R

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

1. Systemic and intestinal antibody titres were measured in chickens following subcutaneous, intraperitoneal (i.p.), oral (p.o.) and combined i.p./p.o. administration of antigen, in soluble, emulsified or microparticulate form. Antigens tested included keyhole limpet haemocyanin (KLH), killed **Campylobacter jejuni** whole cells and purified **campylobacter** flagellin protein. 2. The effect of immunisation with purified flagellin protein or with killed *C. jejuni* whole cells in reducing intestinal colonisation was assessed. The ability of newlyhatched chicks to respond to immunisation was limited, possibly because of the immaturity of the immune system rather than maternal suppression of an immune response. Only 5 to 13 birds that were first immunised when 1-d-old with KLH showed a systemic response, even after 4 immunisations, whereas 10 of 11 birds that were first immunised at 24 d-old responded systemically. 3. In an immunisation and challenge experiment, birds that were immunised twice intraperitoneally, at 16 and 29 d-old, with killed *C. jejuni* whole cells, had fewer *C. jejuni* in the caecal contents than unimmunised control birds. This reduction in intestinal colonisation, to less than 2% of bacterial numbers in control birds, was associated with an increase in specific IgG in intestinal secretions. There was no significant increase in specific IgA or IgM in intestinal secretions following immunisation and challenge. 4. These results indicate that immunisation can reduce the level of intestinal infection with *C. jejuni*. The protection may be enhanced by developing improved methods of immunisation that stimulate production of increased titres of specific antibody in intestinal secretions, particularly specific IgA antibody.

Tags: Animal; Female; Support, Non-U.S. Gov't

Descriptors: Antibodies, Bacterial--metabolism--ME; * **Campylobacter** Infections--veterinary--VE; * **Campylobacter jejuni** --isolation and purification--IP; *Chickens--immunology--IM; *Intestines--microbiology--MI; *Poultry Diseases--immunology--IM; *Poultry Diseases --prevention and control--PC; *Vaccines, Inactivated--pharmacology--PD; Antibodies, Bacterial--immunology--IM; Antigens--immunology--IM; Antigens--metabolism--ME; Antigens, Bacterial--immunology--IM; Antigens, Bacterial--metabolism--ME; **Campylobacter** Infections--immunology--IM; **Campylobacter** Infections--prevention and control--PC; **Campylobacter jejuni**--immunology--IM; Feces--microbiology--MI; Flagellin--immunology--IM; Flagellin--pharmacology--PD; Hemocyanin--immunology--IM; Hemocyanin--pharmacology--PD; Immunoglobulin A--immunology--IM; Immunoglobulin A--metabolism--ME; Immunoglobulin G--immunology--IM; Immunoglobulin G--metabolism--ME; Intestines--drug effects--DE; Time Factors; Vaccines, Inactivated--immunology--IM

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens); 0 (Antigens, Bacterial); 0 (Immunoglobulin A); 0 (Immunoglobulin G); 0 (Vaccines, Inactivated); 12777-81-0 (Flagellin); 9013-72-3 (Hemocyanin)

Record Date Created: 19970206

Record Date Completed: 19970206

10291007 96092877 PMID: 8590089

In ovo oral vaccination with Campylobacter jejuni establishes early development of intestinal immunity in chickens.

Noor S M; Husband A J; Widders P R

Department of Veterinary Pathology, University of Sydney, New South Wales, Australia.

British poultry science (ENGLAND) Sep 1995, 36 (4) p563-73, ISSN 0007-1668 Journal Code: 15740290R

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

1. Chick embryos were orally immunised at day 16 of incubation by injection of heat-killed **Campylobacter jejuni** organisms into the amniotic fluid. The response to vaccination was observed at 5 d after hatching or, in some birds which received a postnatal oral booster vaccination, at 7 d after hatching, and the response was observed at 14 d of age. 2. The titres of antibody in serum, bile and intestinal scrapings, the distribution of immunoglobulin-containing cells in the spleen, duodenum and ileum and the expression on peripheral blood leukocytes (PBL) of the T cell surface markers CD3, CD4 and CD8 were determined. 3. Whereas low titres of anti-flagellin antibody were detected in serum, bile and intestinal scrapings of unimmunised birds, high titres were observed in immunised birds. 4. An increase in antibody of all isotypes was detectable in serum but the elevation in IgA antibody in intestinal scrapings and bile was particularly striking. This response was reflected in a dramatic increase in immunoglobulin-containing cells, detected by fluorescent histology, particularly those associated with IgA and IgM isotypes in the spleen and intestine of immunised birds. 5. Secondary oral boosting after hatching resulted in a depression in serum anti-flagellin antibody in immunised birds compared to pre-boosting titres (although still significantly higher than in non-immunised controls) but an increase in IgA antibody in intestinal scrapings and bile. The number of immunoglobulin-containing cells was also increased after boosting. 6. Neither immunisation regimen caused a significant change in the numbers of circulating CD3, CD4 or CD8 T cells. 7. These results indicate that in ovo oral immunisation with C. jejuni antigens stimulates the precocious development of immunity in chicks.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: **Campylobacter** Infections--veterinary--VE; * **Campylobacter jejuni**--immunology--IM; *Chickens--immunology--IM; *Intestines--immunology--IM; *Poultry Diseases--immunology--IM; *Vaccination--veterinary--VE; Administration, Oral; Amnion--immunology--IM; Antibodies, Bacterial--analysis--AN; Antibodies, Bacterial--blood--BL; Antigens, CD3--analysis--AN; Antigens, CD4--analysis--AN; Antigens, CD8--analysis--AN; Bile--immunology--IM; **Campylobacter** Infections--immunology--IM; **Campylobacter** Infections--prevention and control--PC; Chickens--metabolism--ME; Chickens--microbiology--MI; Immunity, Mucosal; Immunization, Secondary--veterinary--VE; Immunoglobulins--analysis--AN; Immunoglobulins--blood--BL; Intestines--microbiology--MI; Lymphocytes--cytology--CY; Lymphocytes--immunology--IM; Poultry Diseases--prevention and control--PC; Vaccination--methods--MT; Weight Gain--physiology--PH

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens, CD3); 0 (Antigens, CD4); 0 (Antigens, CD8); 0 (Immunoglobulins)

Record Date Created: 19960325

Record Date Completed: 19960325

08383145 95071121 PMID: 7526839

Isotype, specificity, and kinetics of systemic and mucosal antibodies to Campylobacter jejuni antigens, including flagellin, during experimental oral infections of chickens .

Cawthraw S; Ayling R; Nuijten P; Wassenaar T; Newell D G

Central Veterinary Laboratory (Weybridge), New Haw, Addlestone, Surrey, United Kingdom.

Avian diseases (UNITED STATES) Apr-Jun 1994, 38 (2) p341-9, ISSN 0005-2086 Journal Code: 0370617

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

The immune response of chickens to *Campylobacter jejuni* infection was studied as a step in the search for vaccine candidates. One-day-old chicks orally challenged with *C. jejuni* strain 81116 showed significant increases in specific IgG, IgA, and IgM circulating antibodies, as detected by enzyme-linked immunosorbent assay (ELISA). These levels peaked at 9, 5, and 7 weeks postinfection, respectively. Maternal IgG antibodies were also detected over the first 2 weeks. Specific mucosal IgG and IgA antibody levels also increased significantly. All of the birds demonstrated a major response to the 62- kDa flagellin protein by Western blotting techniques. The immunodominance of flagellin was confirmed by ELISA using an antigen preparation from an aflagellate mutant. When overlapping recombinant polypeptide fragments of flagellin were used, epitopes detected by chicken antibodies were observed in region IV, between residues 95-340 of the protein. Thus flagellin may be suitable candidate for a vaccine, although its role in protection must first be established.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: Antibodies, Bacterial--biosynthesis--BI; *Antigens, Bacterial--immunology--IM; * Campylobacter Infections--immunology--IM; * Campylobacter jejuni--immunology--IM; *Flagellin--immunology--IM; * Immunoglobulin G--biosynthesis--BI; * Immunoglobulin Isotypes --biosynthesis--BI; *Intestinal Mucosa--immunology--IM; Antibodies, Bacterial--blood--BL; Antibodies, Bacterial--classification--CL; Antibody Specificity; Blotting, Western; Chickens; Enzyme-Linked Immunosorbent Assay; Epitopes--analysis--AN; Immunoglobulin G--blood--BL; Immunoglobulin G--classification--CL; Immunoglobulin Isotypes --classification--CL

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens, Bacterial); 0 (Epitopes); 0 (Immunoglobulin G); 0 (Immunoglobulin Isotypes); 12777-81-0 (Flagellin)

Record Date Created: 19941129

Record Date Completed: 19941129

11197973 98074583 PMID: 9413103

Campylobacter jejuni in broiler chickens: colonization and humoral immunity following oral vaccination and experimental infection.

Rice B E; Rollins D M; Mallinson E T; Carr L; Joseph S W

Enteric Diseases Program, Naval Medical Research Institute, Bethesda, MD 20889-5607, USA.

Vaccine (ENGLAND) Dec 1997, 15 (17-18) p1922-32, ISSN 0264-410X

Journal Code: 8406899

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

A formalin inactivated, **Campylobacter jejuni** whole cell vaccine, either with or without *Escherichia coli* heat labile toxin (LT) as a mucosal adjuvant, was administered orally to broiler chickens. Three vaccine trials were performed, differing in the number of vaccinations, and time of administration, as well as the inclusion and dose of LT. The overall reductions of *C. jejuni* colonization in the vaccinated chickens ranged from 16 to 93% compared with non-vaccinated controls. Enhanced levels of anti-*C. jejuni* secretory **IgA antibodies** were demonstrated in vaccinated **chickens**. Vaccination also appeared to induce an anamnestic response to *C. jejuni* antigens in the 14-33 kDa range, as demonstrated by Western immunoblots. Interestingly, the inclusion of LT in the vaccine regimen did not appear to boost the immunogenicity of the vaccine. These results are encouraging and suggest that future development of successful oral vaccines for the control of enteropathogenic **Campylobacter** in poultry is feasible.

Tags: Animal; Support, Non-U.S. Gov't; Support, U.S. Gov't, Non-P.H.S.

Descriptors: Bacterial Vaccines--therapeutic use--TU; * **Campylobacter** Infections--veterinary--VE; * **Campylobacter jejuni**--immunology--IM; *Poultry Diseases--immunology--IM; *Poultry Diseases--prevention and control--PC; Administration, Oral; Antibody Formation--immunology--IM; **Campylobacter** Infections--immunology--IM; **Campylobacter** Infections--prevention and control--PC; Poultry Diseases--metabolism--ME; Vaccines, Inactivated--therapeutic use--TU

CAS Registry No.: 0 (Bacterial Vaccines); 0 (Vaccines, Inactivated)

Record Date Created: 19980223

Record Date Completed: 19980223

10257830 96059438 PMID: 7483909

Development of humoral precipitating antibodies to *Campylobacter* spp. in chickens]

Entwicklung humoraler präzipitierender Antikörper gegen *Campylobacter* spp. beim Huhn.

Glunder G

Klinik für Geflügel, Tierärztlichen Hochschule Hannover.

Zentralblatt für Veterinärmedizin. Reihe B. Journal of veterinary medicine. Series B (GERMANY) Apr 1995, 42 (2) p89-99, ISSN 0514-7166
Journal Code: 0331325

Document type: Journal Article ; English Abstract

Languages: GERMAN

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

Development of humoral precipitating antibodies against *Campylobacter* spp. in chickens . The development of precipitating antibodies in chickens was examined by two-dimensional immunodiffusion test after immunization with a formal inactivated vaccine and after subcutaneous and oral application of different live *campylobacter* serovars. The supernatant of bacterial cells after sonication and centrifugation was used as an antigen in the agar-gel precipitin test. Antisera against different *campylobacter* serovars showed a high percentage of cross-reactions. In chickens immunized with an inactivated vaccine at an age of 1, 2, 3, 4 and 7 weeks, precipitating antibodies could be demonstrated for the first time at 7 days p.i. Except for 1-week-old birds, sera from the other groups reacted positively at 14 days p.o. After subcutaneous duplication of live organisms to 4-week-old chickens , antibodies could already be demonstrated at 4 days p.i. later in part of the experimental groups. No interrelation could be detected between antibody titers, measured by enzyme-linked immunoabsorbent assay (ELISA), from precipitating sera, as well as from those from non-precipitating sera. Precipitating antibodies and antibody titers in the ELISA were examined in sera from groups of birds infected at an age of 1, 2, 3, 4 and 7 weeks. During the *Campylobacter* excretion period, a distinct peak of antibody titers occurred in 1- and 7-week old birds, whereas other groups showed a steady increase in titers. Precipitating antibodies were only found in 1- and 2-week-old chickens.

Tags: Animal

Descriptors: Antibodies, Bacterial--biosynthesis--BI; *Bacterial Vaccines--immunology--IM; * *Campylobacter* --immunology--IM; * *Campylobacter* Infections--veterinary--VE; *Chickens; *Poultry Diseases--immunology--IM; *Campylobacter* Infections--immunology--IM; Immunodiffusion--veterinary--VE; Vaccines, Inactivated--immunology--IM

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Bacterial Vaccines); 0 (Vaccines, Inactivated)

Record Date Created: 19951127

Record Date Completed: 19951127

10200549 96001607 PMID: 8526019

Biological properties of yolk immunoglobulins.

Janson A K; Smith C I; Hammarstrom L

Dept of Clinical Immunology, Karolinska Institute, Huddinge Hospital, Sweden.

Advances in experimental medicine and biology (UNITED STATES) 1995, 371A p685-90, ISSN 0065-2598 Journal Code: 0121103

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

Tags: Animal; Comparative Study; Female; Human

Descriptors: Antibodies, Bacterial--immunology--IM; * **Campylobacter jejuni**--immunology--IM; *Egg Proteins--immunology--IM; *Immunoglobulins--immunology--IM; *Shigella flexneri--immunology--IM; Antibodies, Bacterial--isolation and purification--IP; Bacterial Vaccines--immunology--IM; Chickens; Egg Proteins--isolation and purification--IP; Enzyme-Linked Immunosorbent Assay; Granulocytes--immunology--IM; Granulocytes--microbiology--MI; Hela Cells; Hemagglutination Tests; Immunization, Secondary; Immunoglobulins--isolation and purification--IP; Phagocytosis; Shigella flexneri--physiology--PH; Vaccination

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Bacterial Vaccines); 0 (Egg Proteins); 0 (IgY); 0 (Immunoglobulins)

Record Date Created: 19960124

Record Date Completed: 19960124

06814168 91054093 PMID: 2241686

Influence of antibody treatment of Campylobacter jejuni on the dose required to colonize chicks.

Stern N J; Meinersmann R J; Dickerson H W

Poultry Microbiological Safety Research Unit, Richard B. Russell Agricultural Research Center, USDA-Agricultural Research Service, Athens, Georgia 30613.

Avian diseases (UNITED STATES) Jul-Sep 1990, 34 (3) p595-601, ISSN 0005-2086 Journal Code: 0370617

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

This study was designed to clarify the role of antibodies in controlling chicken colonization by *Campylobacter jejuni*. Cecal colonization by *C. jejuni* was compared after the organism was exposed either to phosphate-buffered saline, normal rabbit serum, rabbit hyperimmune anti-*C. jejuni* serum, or anti-*C. jejuni* antibodies extracted from chicken bile. Antibodies from chicken bile were extracted by affinity absorption against outer-membrane proteins from the challenge organism. Sera were heated 1 hour at 56 C to destroy complement activity. Bacterial inoculum levels were enumerated after 1 hour exposure at 4 C to the various treatments. The heated sera and the bile antibodies were not bactericidal, and bacterial agglutination was not evident. Serial dilutions of the antibody-treated *C. jejuni* were given by gavage into 1-day-old chicks. Six days later, the ceca were removed from the chicks, and samples were cultured on *Campylobacter*-charcoal differential agar. The colonization dose-50% was increased by twofold to 160-fold when the organism was preincubated with hyperimmune antiserum or the bile antibodies as compared with preincubation with phosphate-buffered saline. We conclude that antibodies inhibit chicken cecal colonization by *C. jejuni*.

Tags: Animal; Support, Non-U.S. Gov't; Support, U.S. Gov't, Non-P.H.S.

Descriptors: *Campylobacter* Infections--veterinary--VE; * *Campylobacter jejuni*--immunology--IM; *Chickens; *Immunization, Passive; *Poultry Diseases--immunology--IM; Bile--immunology--IM; *Campylobacter* Infections--immunology--IM; *Campylobacter jejuni*--growth and development--GD; Carrier State--immunology--IM; Carrier State--veterinary--VE; Cecum--microbiology--MI; Colony Count, Microbial; Dose-Response Relationship, Immunologic; Enzyme-Linked Immunosorbent Assay; Immune Sera--immunology--IM; Immunoglobulin A, Secretory--immunology--IM; Immunoglobulin A, Secretory--isolation and purification--IP

CAS Registry No.: 0 (Immune Sera); 0 (Immunoglobulin A, Secretory)

Record Date Created: 19901207

Record Date Completed: 19901207

06814167 91054092 PMID: 2241685

Influence of Campylobacter jejuni cecal colonization on immunoglobulin response in chickens .

Myszewski M A; Stern N J

Department of Medical Microbiology, University of Georgia, Athens 30606.

Avian diseases (UNITED STATES) Jul-Sep 1990, 34 (3) p588-94, ISSN 0005-2086 Journal Code: 0370617 .

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

The immunoglobulin response of chickens to colonization by **Campylobacter jejuni** isolates B-540 and Clin-1 was monitored. **Chicken** humoral **IgG** and biliary secretory **IgA** (sIgA) responses were assessed by enzyme-linked immunosorbent assay (ELISA). Samples were taken from 128 C. jejuni-colonized chickens and 104 uncolonized chickens housed in a controlled environment. An indirect ELISA was performed using the homologous isolate of C. jejuni as the capture antigen and was developed with the specific goat anti-chicken **IgG** or **IgA** alkaline phosphatase conjugates. The ELISA absorbance values of the test samples at 405 nm (serum diluted 1:32 and bile diluted 1:10) were normalized in direct proportion to standard sera and bile sample values. In the colonized chickens, humoral **IgG** activities were highest at hatch, dropped to their lowest level after 2 weeks, and increased by 8 weeks to levels similar to those detected at hatch. The sIgA activity was lowest at hatch and increased by 4 weeks in colonized chickens while remaining lower in the control chickens. Chickens colonized with isolate B-540 showed a primary sIgA response during the first 4 weeks and reached a plateau over the final 4 weeks. In spite of these limited humoral and secretory immunoglobulin responses, once the chicken ceca was colonized by C. jejuni, the organism persisted throughout the 8-week experiment.

Tags: Animal

Descriptors: **Campylobacter jejuni**--immunology--IM; *Cecum--microbiology--MI; * **Chickens** --immunology--IM; * **Immunoglobulin A**, Secretory --biosynthesis--BI; *Immunoglobulin G--biosynthesis--BI; Bile--immunology--IM; **Campylobacter** Infections--immunology--IM; **Campylobacter** Infections--veterinary--VE; **Campylobacter jejuni**--growth and development--GD; Carrier State--immunology--IM; Carrier State--veterinary--VE; Enzyme-Linked Immunosorbent Assay; Poultry Diseases--immunology--IM

CAS Registry No.: 0 (Immunoglobulin A, Secretory); 0 (Immunoglobulin G)

Record Date Created: 19901207

Record Date Completed: 19901207

08383145 95071121 PMID: 7526839

Isotype, specificity, and kinetics of systemic and mucosal antibodies to Campylobacter jejuni antigens, including flagellin, during experimental oral infections of chickens.

Cawthraw S; Ayling R; Nuijten P; Wassenaar T; Newell D G
Central Veterinary Laboratory (Weybridge), New Haw, Addlestone, Surrey,
United Kingdom.

Avian diseases (UNITED STATES) Apr-Jun 1994, 38 (2) p341-9, ISSN
0005-2086 Journal Code: 0370617

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

The immune response of chickens to *Campylobacter jejuni* infection was studied as a step in the search for vaccine candidates. One-day-old chicks orally challenged with C. jejuni strain 81116 showed significant increases in specific IgG, IgA, and IgM circulating antibodies, as detected by enzyme-linked immunosorbent assay (ELISA). These levels peaked at 9, 5, and 7 weeks postinfection, respectively. Maternal IgG antibodies were also detected over the first 2 weeks. Specific mucosal IgG and IgA antibody levels also increased significantly. All of the birds demonstrated a major response to the 62- kDa flagellin protein by Western blotting techniques. The immunodominance of flagellin was confirmed by ELISA using an antigen preparation from an aflagellate mutant. When overlapping recombinant polypeptide fragments of flagellin were used, epitopes detected by chicken antibodies were observed in region IV, between residues 95-340 of the protein. Thus flagellin may be suitable candidate for a vaccine, although its role in protection must first be established.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: Antibodies, Bacterial--biosynthesis--BI; *Antigens, Bacterial--immunology--IM; * *Campylobacter* Infections--immunology--IM; * *Campylobacter jejuni*--immunology--IM; *Flagellin--immunology--IM; *Immunoglobulin G--biosynthesis--BI; *Immunoglobulin Isotypes--biosynthesis--BI; *Intestinal Mucosa--immunology--IM; Antibodies, Bacterial--blood--BL; Antibodies, Bacterial--classification--CL; Antibody Specificity; Blotting, Western; Chickens; Enzyme-Linked Immunosorbent Assay; Epitopes--analysis--AN; Immunoglobulin G--blood--BL; Immunoglobulin G--classification--CL; Immunoglobulin Isotypes--classification--CL

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens, Bacterial); 0 (Epitopes); 0 (Immunoglobulin G); 0 (Immunoglobulin Isotypes); 12777-81-0 (Flagellin)

Record Date Created: 19941129

Record Date Completed: 19941129

11732157 99169031 PMID: 10069862

Apoptotic effect of outer-membrane proteins from Campylobacter jejuni on chicken lymphocytes.

Zhu J; Meinersmann R J; Hiatt K L; Evans D L

Poultry Microbiological Safety Research Unit, Russell Agricultural Research Center, U.S. Department of Agriculture, Agricultural Research Service, Athens, GA 30604.

Current microbiology (UNITED STATES) Apr 1999, 38 (4) p244-9, ISSN 0343-8651 Journal Code: 7808448

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS; AIDS/HIV

Campylobacter jejuni is a significant cause of food-borne diseases in humans. The bacterium is considered a commensal organism in **chickens**, and it can heavily colonize **chickens** without causing inflammation. Poultry may be the major reservoir for the human infection in developed countries. Here we show that an outer-membrane protein extract prepared from the bacteria caused apoptosis of **chicken** lymphocytes detected in vitro with the terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling assay that preferentially labels individual apoptotic cells. Blood- and spleen-lymphocytes from different-aged **chickens** displayed a significantly greater percentage of apoptotic cells after culture with the outer-membrane proteins from *C. jejuni* than controls treated with phosphate-buffered saline, **chicken** ovalbumin, or outer-membrane proteins prepared from *E. coli* strain BL21. The *C. jejuni* extract also produced apoptosis of **chicken** lymphoblastoid tumor cell lines. Apoptosis was blocked by pretreating the extract with proteinase K or antiserum against outer-membrane proteins. The results suggest that *C. jejuni* may be capable of achieving **immune** avoidance in **chickens** by causing apoptosis of lymphocytes.

Tags: Animal; Support, U.S. Gov't, Non-P.H.S.

Descriptors: Apoptosis; *Bacterial Outer Membrane Proteins--pharmacology--PD; *CD4-Positive T-Lymphocytes--drug effects--DE; *CD8-Positive T-Lymphocytes--drug effects--DE; * **Campylobacter jejuni**--chemistry--CH; Age Factors; Bacterial Outer Membrane Proteins--drug effects--DE; **Campylobacter jejuni**--immunology--IM; Chickens; Endopeptidase K--pharmacology--PD; *Escherichia coli*--chemistry--CH; Ovalbumin--pharmacology--PD; Specific Pathogen-Free Organisms; Tumor Cells, Cultured CAS Registry No.: 0 (Bacterial Outer Membrane Proteins); 9006-59-1 (Ovalbumin)

Enzyme No.: EC 3.4.21.64 (Endopeptidase K)

Record Date Created: 19990503

Record Date Completed: 19990503

10010 Comparative Biochemistry, General
 10062 Biochemical Studies-Nucleic Acids, Purines and Pyrimidines
 10064 Biochemical Studies-Proteins, Peptides and Amino Acids
 10506 Biophysics-Molecular Properties and Macromolecules
 10508 Biophysics-Membrane Phenomena
 12504 Pathology, General and Miscellaneous-Diagnostic
 15002 Blood, Blood-Forming Organs and Body Fluids-Blood and Lymph
 Studies
 34502 Immunology and Immunochemistry-General; Methods
 36002 Medical and Clinical Microbiology-Bacteriology
 38006 Veterinary Science-Microbiology
 10066 Biochemical Studies-Lipids
 10068 Biochemical Studies-Carbohydrates
 36504 Medical and Clinical Microbiology-Serodiagnosis
 BIOSYSTEMATIC CODES:
 06210 Aerobic Helical or Vibrioid Gram-Negatives (1992-)
 06703 Pasteurellaceae (1992-)
 85536 Galliformes
 85570 Trogoniformes
 85715 Bovidae
 85765 Canidae
 86215 Hominidae

4/9/64 (Item 5 from file: 5)
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09363587 BIOSIS NO.: 199497371957

Isotype, specificity, and kinetics of systemic and mucosal antibodies to
Campylobacter jejuni antigens, including flagellin, during experimental
oral infections of chickens.

AUTHOR: Cawthraw S; Ayling R; Nuijten P; Wassenaar T; Newell D G(a)

AUTHOR ADDRESS: (a)Central Vet. Lab., New Haw, Addlestone, Surrey KT15 3NB

**UK

JOURNAL: Avian Diseases 38 (2):p341-349 1994

ISSN: 0005-2086

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

SUMMARY LANGUAGE: English; Spanish

ABSTRACT: The **immune** response of **chickens** to **Campylobacter jejuni**
 infection was studied as a step in the search for vaccine candidates.
 One-day-old chicks orally challenged with C. jejuni strain 81116 showed
 significant increases in specific IgG, IgA, and IgM circulating
antibodies, as detected by enzyme-linked immunosorbent assay
 (ELISA). These levels peaked at 9, 5, and 7 weeks postinfection,
 respectively. Maternal IgG **antibodies** were also detected over the first
 2 weeks. Specific mucosal IgG and IgA **antibody** levels also increased
 significantly. All of the **birds** demonstrated a major response to the
 62- **kDa** flagellin protein by Western blotting techniques. The
 immunodominance of flagellin was confirmed by ELISA using an antigen
 preparation from an **aflagellate** mutant. When overlapping recombinant
 polypeptide fragments of flagellin were used, epitopes detected by
chicken antibodies were observed in region IV, between residues 95-340
 of the protein. Thus flagellin may be a suitable candidate for a vaccine,
 although its role in protection must first be established.

DESCRIPTORS:

MAJOR CONCEPTS: Dental and Oral System (Ingestion and Assimilation);
 Immune System (Chemical Coordination and Homeostasis); Infection;
 Pathology; Veterinary Medicine (Medical Sciences)

BIOSYSTEMATIC NAMES: Aerobic Helical or Vibrioid Gram-Negatives--
 Eubacteria, Bacteria; Galliformes--Aves, Vertebrata, Chordata, Animalia

ORGANISMS: aerobic helical or vibrioid gram-negative bacteria (Aerobic
 Helical or Vibrioid Gram-Negatives); **Campylobacter jejuni** (Aerobic

Helical or Vibrioid Gram-Negatives); Galliformes (Galliformes
 BIOSYSTEMATIC CLASSIFICATION (SUPER TAXA): animals; bacteria; birds;

chordates; eubacteria; microorganisms; nonhuman vertebrates;
vertebrates

MISCELLANEOUS TERMS: PATHOGENICITY; VACCINATION

CONCEPT CODES:

12512 Pathology, General and Miscellaneous-Therapy (1971-)
19006 Dental and Oral Biology-Pathology
34504 Immunology and Immunochemistry-Bacterial, Viral and Fungal
36002 Medical and Clinical Microbiology-Bacteriology
38004 Veterinary Science-Pathology
38006 Veterinary Science-Microbiology
10064 Biochemical Studies-Proteins, Peptides and Amino Acids
10068 Biochemical Studies-Carbohydrates

BIOSYSTEMATIC CODES:

06210 Aerobic Helical or Vibrioid Gram-Negatives (1992-)
85536 Galliformes

4/9/68 (Item 3 from file: 10)

DIALOG(R)File 10:AGRICOLA

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Isotype, specificity, and kinetics of systemic and mucosal antibodies to Campylobacter jejuni antigens, including flagellin, during experimental oral infections of chickens

Cawthraw, S. Ayling, R.; Nuijten, P.; Wassenaar, T.; Newell, D.G.
Kennett Square, Pa. : American Association of Avian Pathologists Inc.
Avian diseases. Apr/June 1994. v. 38 (2) p. 341-349.

ISSN: 0005-2086 CODEN: AVDIAI

DNAL CALL NO: 41.8 Av5

Language: English Summary Language: Spanish

Includes references

Place of Publication: Pennsylvania

Subfile: IND; OTHER US (NOT EXP STN, EXT, USDA; SINCE 12/76);

Document Type: Article

The **immune** response of **chickens** to **Campylobacter jejuni** infection was studied as a step in the search for vaccine candidates. One-day-old chicks orally challenged with C. jejuni strain 81116 showed significant increases in specific IgG, IgA, and IgM circulating **antibodies**, as detected by enzyme-linked immunosorbent assay (ELISA). These levels peaked at 9, 5, and 7 weeks postinfection, respectively. Maternal IgG **antibodies** were also detected over the first 2 weeks. Specific mucosal IgG and IgA **antibody** levels also increased significantly. All of the **birds** demonstrated a major response to the 62- kDa flagellin protein by Western blotting techniques. The immunodominance of flagellin was confirmed by ELISA using an antigen preparation from an **aflagellate** mutant. When overlapping recombinant polypeptide fragments of flagellin were used, epitopes detected by **chicken antibodies** were observed in region IV, between residues 95-340 of the protein. Thus flagellin may be a suitable candidate for a vaccine, although its role in protection must first be established.

DESCRIPTORS: chickens; **campylobacter jejuni**; experimental infections; antibody formation; iga; igg; igm; maternal antibodies; flagella; bacterial antigens; antigenic determinants; isotypes;

Section Headings: L832 ANIMAL DISEASES-BACTERIAL; L810 VETERINARY PHARMACOLOGY AND IMMUNE THERAPEUTIC AGENTS

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J. Bacteriol., Jun 1992, 3874-3883, Vol 174, No. 12
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Biochemical and antigenic properties of the *Campylobacter* flagellar hook protein

ME Power, RA Alm and TJ Trust

Department of Biochemistry and Microbiology, University of Victoria, British Columbia, Canada.

The flagellar filament-hook complex was removed from *Campylobacter* cells by shearing and was purified by differential solubilization and ultracentrifugation at pH 11 followed by cesium chloride buoyant density ultracentrifugation. Flagellar filaments were then dissociated in 0.2 M glycine-HCl (pH 2.2), and purified hooks were collected by ultracentrifugation. The hooks (105 by 24 nm) each displayed a conical protrusion at the proximal end, a concave cavity at the distal end, and helically arranged subunits. The apparent subunit molecular weight of the hook protein of seven of the eight *Campylobacter* strains studied was 92,500, while that of the other was 94,000. N-terminal amino acid analysis of the hook protein of two strains of *Campylobacter coli* and one strain of *Campylobacter jejuni* demonstrated that the first 15 residues were identical. Amino acid composition analysis showed that the *Campylobacter* hook protein contained 35.7% hydrophobic and 9.5% basic residues. Isoelectric focusing determined that the hook protein was acidic, with a pI of 4.9. Comparisons with the *Salmonella* and *Caulobacter* hook protein compositions

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Infect. Immun., Oct 1994, 4256-4260, Vol 62, No. 10
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Heat shock- and alkaline pH-induced proteins of *Campylobacter jejuni*: characterization and immunological properties

YL Wu, LH Lee, DM Rollins and WM Ching

Naval Medical Research Institute, Bethesda, Maryland 20889-5607.

The protein response to physiological stress was characterized in *Campylobacter jejuni* 81176 after exposure to heat and pH shock and following periods of recovery. Immunoreactivities of major stress-related proteins were determined with anti-*Campylobacter* immune rabbit serum and intestinal lavage fluid. Distinct proteins with molecular masses ranging from 10 to 120 kDa were induced and/or released by selective heat or pH treatments. The most notable responses were those of two proteins with apparent molecular masses of 45 and 64 kDa that were induced and two other proteins of 10 and 12 kDa that were released by selective heat shock, alkaline pH treatment, or both. On the basis of N-terminal sequence analysis and immunological cross-reactivity data, the 64- and 10-kDa proteins were the *C. jejuni* homologs of *Escherichia coli* GroEL and GroES proteins, respectively. Enhanced chemiluminescence Western blotting (immunoblotting) revealed that all four proteins were among the major protein antigens recognized by anti-

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Entry name	Q9R4E6
Primary accession number	Q9R4E6
Secondary accession numbers	None
Entered in TrEMBL in	Release 13, May 2000
Sequence was last modified in	Release 13, May 2000
Annotations were last modified in	Release 25, September 2003

Name and origin of the protein

Protein name	groEL-like stress protein 62 kDa subunit [Fragment]
Synonyms	None
Gene name	None
From	<u>Campylobacter jejuni</u> [TaxID: 197]
Taxonomy	<u>Bacteria</u> ; <u>Proteobacteria</u> ; <u>Epsilonproteobacteria</u> ; <u>Campylobacterales</u> ; <u>Campylobacteraceae</u> ; <u>Campylobacter</u> .

References

- [1] SEQUENCE.
MEDLINE=96123358; PubMed=8577276; [NCBI, ExPASy, EBI, Israel, Japan]
Takata T., Wai S.N., Takade A., Sawae Y., Ono J., Amako K.;
"The purification of a GroEL-like stress protein from aerobically adapted *Campylobacter jejuni*.";
Microbiol. Immunol. 39:639-645(1995).

Comments

None

Cross-references

0318986 96121222 PMID: 8576327

Identification and characterization of an immunogenic outer membrane protein of *Campylobacter jejuni*.

Burnens A; Stucki U; Nicolet J; Frey J

Institute for Veterinary Bacteriology, University of Berne, Switzerland.

Journal of clinical microbiology (UNITED STATES) Nov 1995, 33 (11)
p2826-32, ISSN 0095-1137 Journal Code: 7505564

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: Completed

Subfile: INDEX MEDICUS

We cloned and expressed in *Escherichia coli* a gene encoding an 18- kDa outer membrane protein (Omp18) from *Campylobacter jejuni* ATCC 29428. The nucleotide sequence of the gene encoding Omp18 was determined, and an open reading frame of 165 amino acids was revealed. The amino acid sequence had the typical features of a leader sequence and a signal peptidase II cleavage site at the N-terminal part of Omp18. Moreover, the sequence had a high degree of similarity to the peptidoglycan-associated outer membrane lipoprotein P6 of *Haemophilus influenzae* and the peptidoglycan-associated lipoprotein PAL of *E. coli*. Southern blot analysis in which the cloned gene was used as a probe revealed genes similar to that encoding Omp18 in all species of the thermophilic group of *campylobacters* as well as *Campylobacter sputorum*. All *campylobacters* tested expressed a protein with a molecular mass identical to that of Omp18. The protein reacted immunologically with **polyclonal antibodies** directed against Omp18 from *C. jejuni*. PCR amplification of the gene encoding Omp18 with specific primers and subsequent restriction enzyme analysis of the amplified DNA fragments showed that the gene for Omp18 is highly conserved in *C. jejuni* strains isolated from humans, dogs, cats, calves, and **chickens** but is different in other *Campylobacter* species. In order to obtain pure recombinant Omp18 protein for serological assays, the cloned gene for Omp18 was genetically modified by replacing the signal sequence with a DNA segment encoding six adjacent histidine residues. Expression of this construct in *E. coli* allowed purification of the modified protein (Omp18-6xHis) by metal chelation chromatography. Sera from patients with past *C. jejuni* infection reacted positively with Omp18-6xHis, while sera from healthy blood donors showed no reaction with this antigen. Omp18, which is an outer membrane protein belonging to the family of PALs is well conserved in *C. jejuni* and is highly immunogenic. It is therefore a good candidate as an antigen for the serological diagnosis of past *C. jejuni* infections.

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[\[Keywords\]](#) [\[Features\]](#) [\[Sequence\]](#) [\[Tools\]](#)

Note: most headings are clickable, even if they don't appear as links. They link to the user manual or other documents.

Entry information

Entry name	CH60_GAMJE
Primary accession number	O69289
Secondary accession number	Q9PN75
Entered in Swiss-Prot in	Release 39, May 2000
Sequence was last modified in	Release 40, October 2001
Annotations were last modified in	Release 41, February 2003
Name and origin of the protein	
Protein name	60 kDa chaperonin
Synonyms	Protein Cpn60 groEL protein
Gene name	GROL or GROEL or MOPA or <u>CJ1221</u>
From	<u>Campylobacter jejuni</u> [TaxID: 197]
Taxonomy	Bacteria; <u>Proteobacteria</u> ; <u>Epsilonproteobacteria</u> ; <u>Campylobacterales</u> ; <u>Campylobacteraceae</u> ; <u>Campylobacter</u> .
References	

[1]	<p>SEQUENCE FROM NUCLEIC ACID.</p> <p>MEDLINE=99140140; PubMed=10206714; [NCBI, ExPASy, EBI, Israel, Japan]</p> <p>Thies F.L., Weishaupt A., Karch H., Hartung H.P., Giegerich G.;</p> <p>"Cloning, sequencing and molecular analysis of the <i>Campylobacter jejuni</i> groESL bicistronic operon.";</p> <p><i>Microbiology</i> 145:89-98(1999).</p>
[2]	<p>SEQUENCE FROM NUCLEIC ACID.</p> <p>STRAIN=ATCC 43429, ATCC 43432, ATCC 43438, and ATCC 43456;</p> <p>Cunningham A., Taboada E., Nash J.H., Wakarchuk W.W., Gilbert M.;</p> <p>"Sequencing of the cpn60 gene from various <i>Campylobacter jejuni</i> isolates.";</p> <p>Submitted (DEC-2001) to the EMBL/GenBank/DDBJ databases.</p>
[3]	<p>SEQUENCE FROM NUCLEIC ACID.</p> <p>STRAIN=NCTC 11168;</p> <p>MEDLINE=20150912; PubMed=10688204; [NCBI, ExPASy, EBI, Israel, Japan]</p> <p>Parkhill J., Wren B.W., Mungall K., Ketley J.M., Churcher C., Basham D., Chillingworth T., Davies R.M., Feltwell T., Holroyd S., Jagels K., Karlyshev A.V., Moule S., Pallen M.J., Penn C.W., Quail M.A., Rajandream M.A., Rutherford K.M., van Vliet A.H.M., Whitehead S., Barrell B.G.;</p> <p>"The genome sequence of the food-borne pathogen <i>Campylobacter jejuni</i> reveals hypervariable sequences.";</p> <p><i>Nature</i> 403:665-668(2000).</p>

Comments

FUNCTION: Prevents misfolding and promotes the refolding and proper assembly of unfolded polypeptides generated under stress conditions (*By similarity*).

SUBUNIT: Oligomer of 14 subunits composed of two stacked rings of 7 subunits (*By similarity*).

SUBCELLULAR LOCATION: Cytoplasmic (*By similarity*).

SIMILARITY: Belongs to the chaperonin (HSP60) family.

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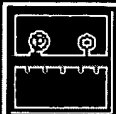
EMBL	Y13334; CAA73778.1; -. [EMBL / GenBank / DDBJ] [CoDingSequence] AF461064; [EMBL / GenBank / DDBJ] AAL76936.1; -. [CoDingSequence] AF461534; [EMBL / GenBank / DDBJ] AAL67841.1; -. [CoDingSequence] AF461535; [EMBL / GenBank / DDBJ] AAL67842.1; -. [CoDingSequence] AF461537; [EMBL / GenBank / DDBJ] AAL67844.1; -. [CoDingSequence] AL139077; [EMBL / GenBank / DDBJ] CAB73475.1; -. [CoDingSequence]
PIR	G81328; G81328.
HSSP	P06139; 1GRL. [HSSP ENTRY / SWISS-3DIMAGE / PDB]
CMR	O69289; CJ1221.
HAMAP	MF_00600; -: 1. PBIL [Family / Alignment / Tree]
InterPro	IPR001844; Chaprnin_Cpn60. IPR002423; Cpn60/TCP-1. Graphical view of domain structure.
Pfam	PF00118; cpn60_TCP1; 1.
PRINTS	PR00298; CHAPERONIN60. PR00304; TCOMPLEXTCP1.
PROSITE	PS00296; CHAPERONINS_CPN60; 1.
ProDom	[Domain structure / List of seq. sharing at least 1 domain]
HOBACGEN	[Family / Alignment / Tree]
BLOCKS	O69289.
ProtoNet	O69289.
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Keywords

Chaperone; ATP-binding; Complete proteome.

Features



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CONFLICT	<u>179</u>	<u>179</u>		A -> P (IN REF. <u>1</u>) .
CONFLICT	<u>383</u>	<u>383</u>		A -> T (IN REF. <u>1</u>) .

Sequence information

Length: 545 AA	Molecular weight: 57970 Da	CRC64: 4DA80BC64330C41E [This is a checksum on the sequence]
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10	20	30	40	50	60
MAKEIIFSDE	ARNKLYEGVK	KLNDAVKVTM	GPRGRNVLIQ	KSFGAPSITK	DGVSVAKEVE
70	80	90	100	110	120
LKDSLENMGA	SLVREVASKT	ADQAGDGTTT	ATVLAHAIFK	EGLRNITAGA	NPIEVKRGMD
130	140	150	160	170	180
KACEAIVAEL	KKLSREVKDK	KEIAQVATIS	ANSDEKIGNL	IADAMEKVGK	DGVITVEEAK
190	200	210	220	230	240
SINDELNVVE	GMQFDRGYLS	PYFITNAEKM	TVELSSPYIL	LFDKKITNLK	DLLPVLEQIQ
250	260	270	280	290	300
KTGKPLLIIA	EDIEGEALAT	LVVNKL RGV L	NISAVKAPGF	GDRRKAMLED	IAILTGGEVI
310	320	330	340	350	360
SEELGRTLES	ATIQDLGQAS	SVIIDKDNTT	IVNGAGEKAN	IDARVNQIKA	QIAETTSDDYD
370	380	390	400	410	420
REKLQERLAK	LSGGVAVIKV	GAATETEMKE	KKDRVDDALS	ATKAAVEEGI	VIGGGAALIK
430	440	450	460	470	480
AKAKIKLDLQ	GDEAIGAAIV	ERALRAPLRQ	IAENAGFDAG	VVNSVENAK	DENTGFDAAK
490	500	510	520	530	540
GEYVNMLES	IIDPVKVERV	ALLNAVSVAS	MLLTTEATIS	EIKEDKPTMP	DMSGMGGMGG
MGGMM					
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